WHAT IS CLAIMED IS:

- 1. A method of creating hybrid proteins having a common biological activity comprising the steps of:
- (a) creating a library comprising 32 or more nucleic acids encoding a plurality of hybrid protein members, wherein the members differ from a set of at least two parent proteins with corresponding amino acids, and
- i. where the parent proteins are homologous proteins having greater than 60% amino acid similarity to each other and having at least one common biological activity,
- ii. where a majority of the library members have a greater than 60% amino acid similarity to any of the parent proteins, and
- iii. where the majority of differences between the library members and the parent proteins are confined to those corresponding amino acids that differ among the parent proteins;
- (b) expressing protein from at least one library member to create at least one hybrid protein;
- (c) selecting at least one protein having a common biological activity of the parent proteins.
 - 2. The method of claim 1, wherein the parent proteins are enzymes.
 - 3. The method of claim 1, wherein the parent proteins are isozymes.
 - 4. The method of claim 1, wherein the parent proteins are polymerases.
- 5. The method of claim 1, wherein the parent proteins have greater than 80% amino acid similarity to each other and the majority of the library members have greater than 80% amino acid similarity to any of the wild-type proteins.
- 6. A library comprising nucleic acids encoding a plurality of hybrid protein members, wherein the members differ from a set of at least two parent proteins with corresponding amino acids, and
- i. where the parent proteins are homologous proteins having greater than 60% amino acid similarity to each other and having at least one common biological activity,

- ii. where a majority of the library members have a greater than 60% amino acid similarity to any of the parent proteins, and
- iii. where the majority of differences between the library members and their parent proteins are confined to those corresponding amino acids that differ among the parent proteins.
 - 7. The method of claim 6, wherein the parent proteins are enzymes.
 - 8. The method of claim 6, wherein the parent proteins are isozymes.
 - 9. The method of claim 6, wherein the parent proteins are polymerases.
- 10. The method of claim 6, wherein the parent proteins have greater than 80% amino acid similarity to each other and the majority of the library members have greater than 80% amino acid similarity to any of the parent proteins.
- 11. A synthetic hybrid protein comprising greater than 60% amino acid similarity to each member of a set of at least two parent proteins, where each parent protein in the set shares greater than 60% amino acid similarity and at least one common biological activity with each member of the set, and wherein the set comprises a subset of invariant amino acids that are identical among all members of the set, and wherein the hybrid protein:
 - (a) shares at least one biological activity with all members of the set;
 - (b) comprises at least 95% of the subset of invariant amino acids
- (c) has a minimum of 5 amino acid residue differences from any member of the set; and
- (d) contains at least 5 variable amino acid residues corresponding to alternating parent proteins.
- 12. The hybrid protein of claim 11, wherein the parent protein is an enzyme.
- 13. The hybrid protein of claim 11, wherein the parent protein is an isozyme.
- 14. The hybrid protein of claim 11, wherein the parent protein is a polymerase.

- 15. The hybrid protein of claim 11, wherein the synthetic protein comprises greater than 80% amino acid similarity to each member of the set and wherein each wild-type protein in the set shares greater than 80% amino acid similarity with each member of the set.
- 16. A hybrid protein of claim 11, wherein the set of parent proteins comprises the *Pyroccus furiosus* family B DNA polymerase and *Pyrococcu ssp.* GB-D DNA Polymerase and the differences from any member of the set comprise at least 10 of the mutations selected from the group listed in Figure 8.